

**S5 Table. SWI/SNF Complex genes.** Burden testing results (based on 1063 MM/MGUS cases and 964 unaffected controls), SGS and prioritized SNV results, and tolerance to missense and loss of function variants (based on ExAC population data).

Gene	Position	Burden (p-value)	SGS	SNV	Intolerance to MS (Z)	Intolerance to LoF (pLI)
<i>ARID1A</i>	1:27022522-27108601	0.001 <sup>‡</sup>	2 pedigrees, borderline suggestive	Y	4.10	1.00
<i>ACTL6A</i>	3:179280708-179306193	0.001 <sup>‡</sup>			2.93	0.99
<i>SMARCC1</i>	3:47627378-47823405	0.821	borderline suggestive		2.39	1.00
<i>PBRM1</i>	3:52579368-52713739	0.505			3.48	1.00
<i>ARID1B</i>	6:157099064-157531913	0.002 <sup>‡</sup>			3.39	1.00
<i>ACTL6B</i>	7:100240726-100254084	0.187			4.04	0.99
<i>SMARCD3</i>	7:150936059-150945749	0.001 <sup>‡</sup>			3.16	0.23
<i>SMARCA2</i>	9:2015342-2193623	0.334			5.57	1.00
<i>ARID2</i>	12:46123620-46301819	0.045 <sup>+</sup>			1.87	1.00
<i>SMARCD1</i>	12:50478983-50494494	0.458			3.95	1.00
<i>SMARCC2</i>	12:56555636-56583351	0.001 <sup>‡</sup>			4.26	1.00
<i>SMARCE1</i>	17:38783976-38804103	0.001 <sup>‡</sup>			2.96	1.00
<i>SMARCD2</i>	17:61909441-61920351	0.685			2.37	0.98
<i>SMARCA4</i>	19:11071598-11172958	0.001 <sup>‡</sup>			8.36	1.00
<i>SMARCB1</i>	22:24129150-24176705	-			4.51	1.00

**Legend:** Position – build HG19; Burden – p-values based on the c-alpha test of high-impact variants with AAF < 0.001 (see Methods section), “-” indicates gene not tested (no variants observed), <sup>‡</sup>significant after multiple testing correction p < 0.0033 (=0.05/15), <sup>+</sup>nominally significant p < 0.05; SGS – gene captured by a border-line suggestive shared genomic segment; SNV – single nucleotide variant with AAF < 0.001, high or moderate deleteriousness, and observed segregating in a high-risk MM pedigree or pathogenic in ClinVar; Intolerance to MS – the gene’s intolerance to missense variants based on analysis of ExAC data<sup>41</sup>, signed Z score based on deviation of observed counts from expected, positive Z indicates intolerance to variation; Intolerance to LoF – based on analysis of ExAC data<sup>41</sup>, Loss of Function (LoF) variants include splice donor or acceptor or non-sense SNVs, genes with a probability of LoF Intolerance (pLI) >= 0.9 are considered extremely intolerant to LoF SNVs.